Amendments to the Claims

This listing of claims replaces all prior listings, and versions, of claims in the application.

Listing of Claims:

- 1 271. (canceled)
- 272. (currently amended) A method of identifying an agent that modulates an eye a retinal abnormality, the method comprising: (a) providing a transgenic mouse whose genome comprises a disruption knockout of the gene which encodes for the native sequence PRO224 polypeptide, said mouse comprising a retinal abnormality resulting from said PRO224 gene knockout; (b) measuring a physiological characteristic of an eye a retina of the transgenic mouse of (a); (c) comparing the measured physiological characteristic of (b) with that of a gender matched wild-type mouse, wherein the physiological characteristic of an eye a retina of the transgenic mouse that differs from the physiological characteristic a retina of the wild-type mouse is identified as an eye a retinal abnormality resulting from the PRO224 gene disruption knockout in the transgenic mouse; (d) administering a test agent to the transgenic mouse of (a); and (e) determining whether the test agent modulates said eye retinal abnormality, whereby an agent which is determined to modulate an eye a retinal abnormality is identified.
- 273. (currently amended) The method of Claim 272, wherein the eye abnormality is a retinal abnormality comprising comprises a retinal artery obstruction or occlusion.
 - 274-279. (canceled)
- 280. (currently amended) The method of Claim 272, wherein the eye <u>retinal</u> abnormality is a <u>vascular</u> retinal abnormality.
- 281. (currently amended) The method of Claim 280 272, wherein the eye abnormality is a retinal abnormality consistent with causes vision problems or blindness.
- 282. (currently amended) The method of Claim 280 272, wherein the retinal abnormality is consistent with retinitis pigmentosa.
- 283. (currently amended) The method of Claim 280 272, wherein the retinal abnormality is characterized by retinal degeneration or retinal dysplasia.
 - 284. (currently amended) The method of Claim 280 272, wherein the retinal abnormality

is consistent with selected from retinal dysplasia, various retinopathics a retinopathy, including retinopathy of prematurity, retrolental fibroplasia, neovascular glaucoma, age-related macular degeneration, diabetic macular edema, corneal neovascularization, corneal graft neovascularization, corneal graft rejection, retinal/choroidal neovascularization, neovascularization of the angle (rubeosis), ocular neovascular disease, vascular restenosis, arteriovenous malformations (AVM), meningioma, hemangioma, angiofibroma, thyroid hyperplasias hyperplasia, (including Grave's disease), corneal and other tissue transplantation, retinal artery obstruction or occlusion; retinal degeneration causing secondary atrophy of the retinal vasculature, retinitis pigmentosa, macular dystrophies, Stargardt's disease, congenital stationary night blindness, choroideremia, gyrate atrophy, Leber's congenital amaurosis, retinoschisis disorders, Wagner's syndrome, Usher syndrome, Zellweger syndrome, Saldino-Mainzer syndrome, Senior-Loken syndrome, Bardet-Biedl syndrome, Alport's syndrome, Alstrom's syndrome, Cockayne's syndrome, dysplasia spondyloepiphysaria congentia, Flynn-Aird syndrome, Friedreich ataxia, Hallgren syndrome, Marshall syndrome, Albers-Schnoberg disease, Refsum's disease, Kearns-Sayre syndrome, Waardenburg's syndrome, Alagile syndrome, myotonic dystrophy, olivopontocerebellar atrophy, Pierre Marie syndrome, Stickler syndrome, carotinemeia, cystinosis, Wolfram syndrome, Bassen-Kornzweig syndrome, abetalipoproteinemia, incontinentia pigmenti, Batten's disease, mucopolysaccharidoses, homocystinuria, or and mannosidosis.

285 - 290. (canceled)

291. (currently amended) The method of Claim 272, wherein the transgenic mouse exhibits the following physiological characteristic an increased mean artery-to-vein ratio associated with retinal degeneration, as compared with gender-matched wild-type littermates: an increased mean artery-to-vein ratio associated with retinal degeneration.

292-386. (canceled)